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(54) Alkali-Soluble Siloxane Polymer, Silmethylene Polymer, and Polyorganosilsesquioxane Polymer.

(57) There are provided an alkali-soluble siloxane polymer, an alkali-soluble silmethylen polymer, and an alkali-soluble polyorganosilsesquioxane polymer. They are useful as a photoresist for the fabrication of semiconductor devices. They are suitable for dry etching because of their superior resistance to oxygen plasmas.

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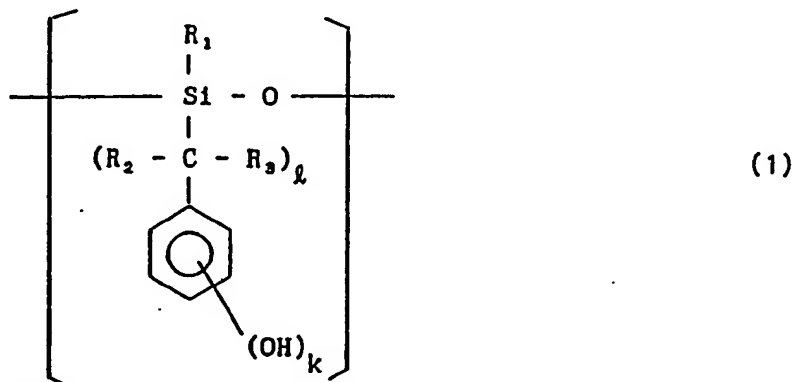
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1 ALKALI-SOLUBLE SILOXANE POLYMER, SILMETHYLENE POLYMER, AND
POLYORGANOSILSESQUIOXANE POLYMER

5 BACKGROUND OF THE INVENTION

The present invention relates to a new polymer which is extremely useful as a functional polymeric material such as light-sensitive and radiation-sensitive materials. More particularly, it relates to an
10 alkali-soluble siloxane polymer in which more than 40% of the siloxane skeleton is composed of the unit represented by the formula (1); an alkali-soluble silmethylen polymer composed of the constitutional unit represented by the formula (2); and an alkali-soluble polyorganosil-
15 sesquioxane polymer represented by the formula (3).



25 (where R_1 is hydrogen or a monovalent organic group other than aromatic groups; R_2 and R_3 are each hydrogen, chlorine, or a monovalent organic group; l is an integer of 1 to 6; and k is 1 or 2.)



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circuits are produced by means of microetching with light or radiation.

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25 polyvinyl phenol and bisazide forms a negative photoresist.

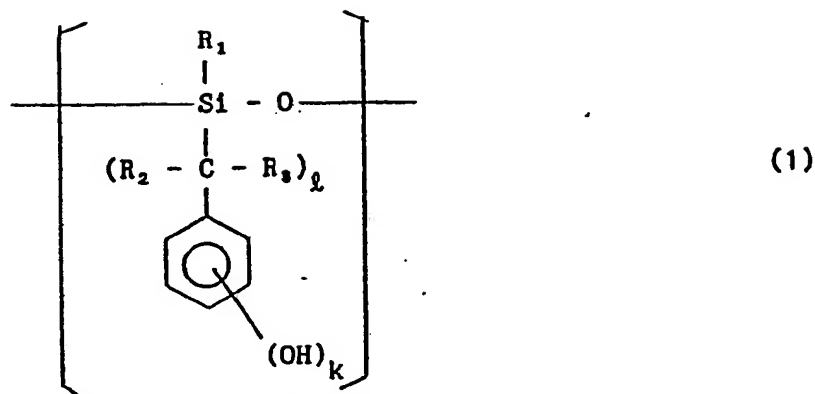
1 After patterning of the resist layer, the exposed substrate under-
goes etching. In view of the trend to ever smaller circuit patterns of
semiconductor devices, wet etching is being replaced by dry etching.
Thus, a resist material is required to withstand dry etching. In the
5 case where the substrate is an organic material, etching is carried out
by an oxygen plasma. The conventional resist material is poor in
resistance to oxygen plasmas, and there is a demand for the improvement
in this regard.

In the meantime, the resist of alkali-development type is described
10 in "Kodak Microelectronics Seminar Proceedings, 116 (1976), by
J. C. Strieter.

SUMMARY OF THE INVENTION

It is an object of the present invention to provide an alkali-
soluble siloxane polymer which is superior in resistance to oxygen
15 plasmas.

The present inventors synthesized a variety of polymers in which
the main chain is a siloxane skeleton and the side chain has phenolic
hydroxyl groups. As the result, it was found that the above-mentioned
object is achieved by a siloxane polymer which contains more than 40% of
20 the unit represented by the following formula (1) in the siloxane
skeleton.

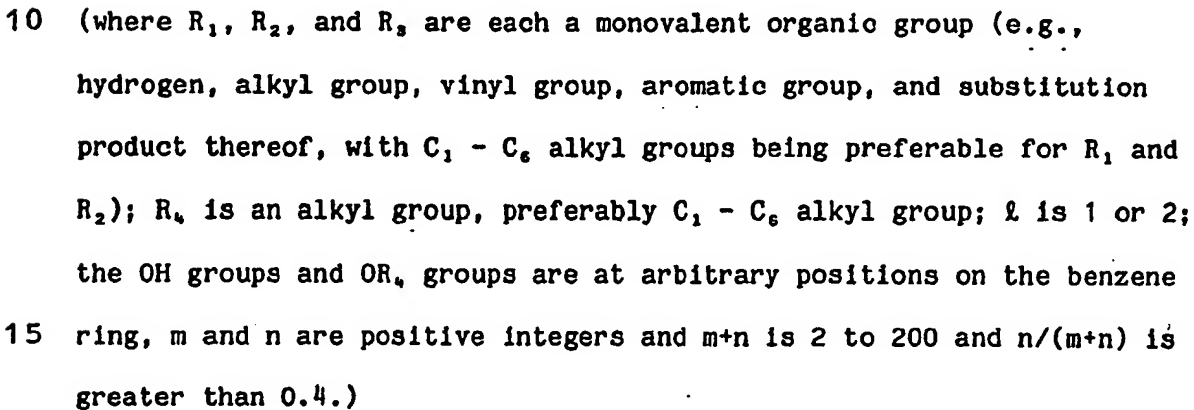


(where R_1 is hydrogen or a monovalent organic group other than aromatic groups (e.g., alkyl groups having 1 to 5 carbon atoms, vinyl group, and substitution product thereof, preferably $-\text{CH}_3$, $-\text{C}_2\text{H}_5$, and $-\text{C}_3\text{H}_7$); R_2 and R_3 are each hydrogen, chlorine, or a monovalent organic group (e.g., alkyl groups having 1 to 5 carbon atoms, vinyl group, aromatic groups such as phenyl and tolyl, and substitution product thereof), with hydrogen being preferable; l is an integer of 1 to 6; and k is 1 or 2. The OH group should preferably be at the p-position of the benzene ring.

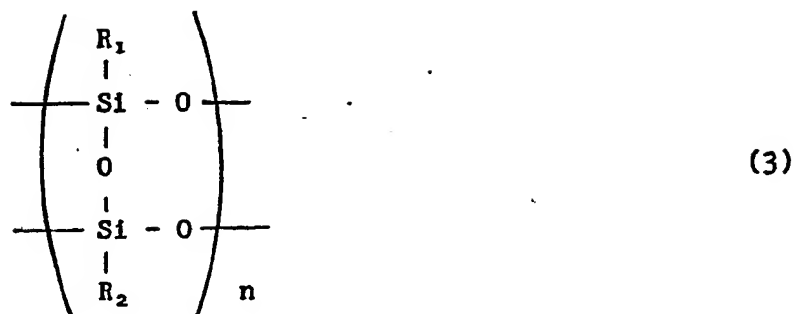
What is important in the present invention is that the aromatic group is not directly connected to silicon in order to avoid the cleavage of the bond between the silicon atom and the aromatic group by the phenolic hydroxyl group. Thus, the siloxane other than that specified by the above formula (1) is not specifically limited unless an aromatic group is directly connected to the silicon atom. The siloxane polymer of this invention will be poor in alkali solubility unless the siloxane skeleton contains more than 40% of the unit represented by the above formula (1).

The present inventors also synthesized a variety of polymers in which the main chain is a silmethylenes skeleton and the side chain has a

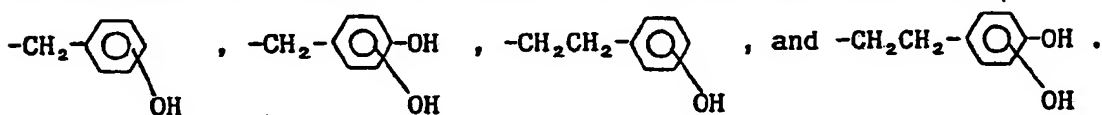
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20 As the result, it was found that the above-mentioned object is achieved
by a polyorganosilsesquioxane represented by the following formula (3).

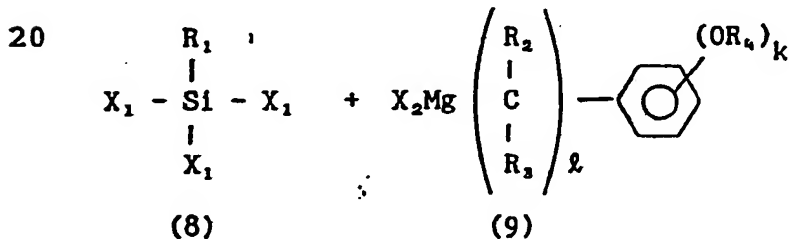


(where R_1 and R_2 are the same or different organic groups; and n is a degree of polymerization.) The polyorganosilsesquioxane may contain more than 40% of R_1 and R_2 each having phenolic hydroxyl groups. The value of n is 8 to 200. Examples of R_1 and R_2 are $C_1 - C_6$ alkyl groups having phenol or cresol as a substituent group, as shown below.

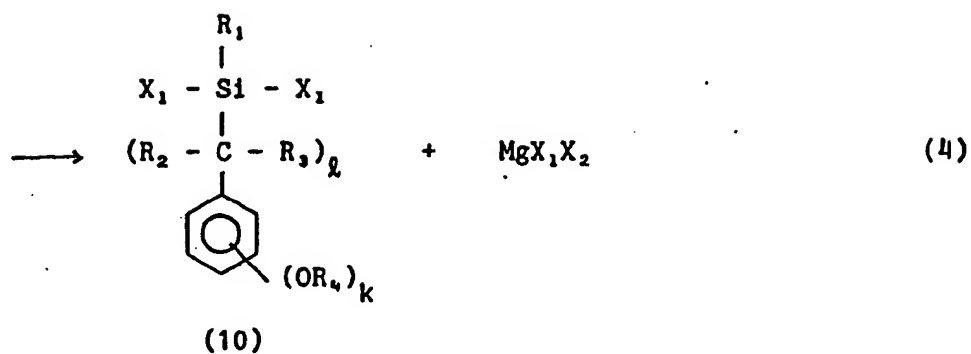


Side chains other than those shown above may be ones in which the phenolic hydroxyl groups are substituted by alkoxy groups, t-butyl-dimethylsiloxy groups, or methylene acetal. This polymer is not sufficiently alkali soluble unless the organic groups having phenolic hydroxyl groups account for more than 40% in the side chains.

The polymer of this invention as represented by the formula (1) is synthesized according to the following equations (4) to (7).

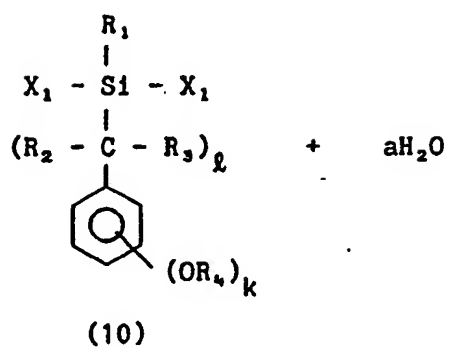


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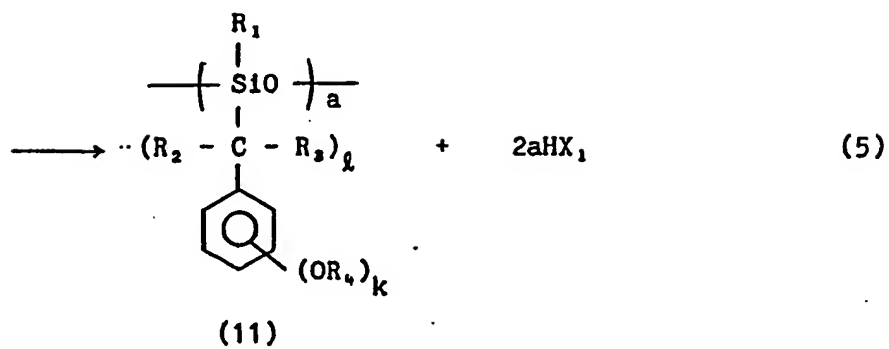


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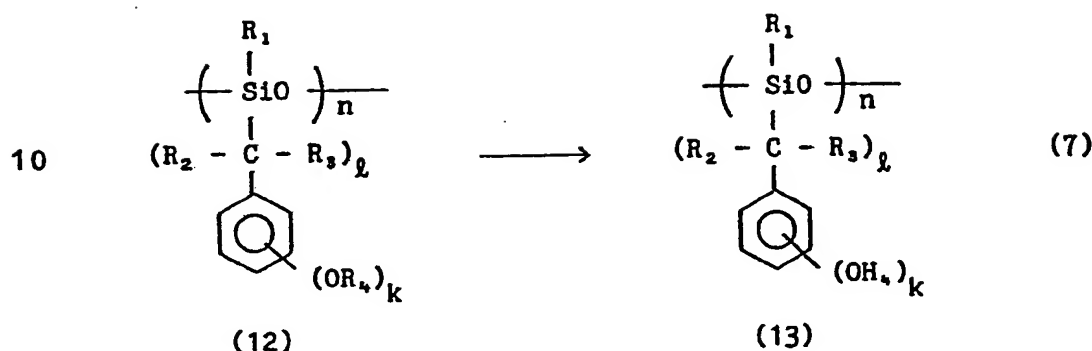
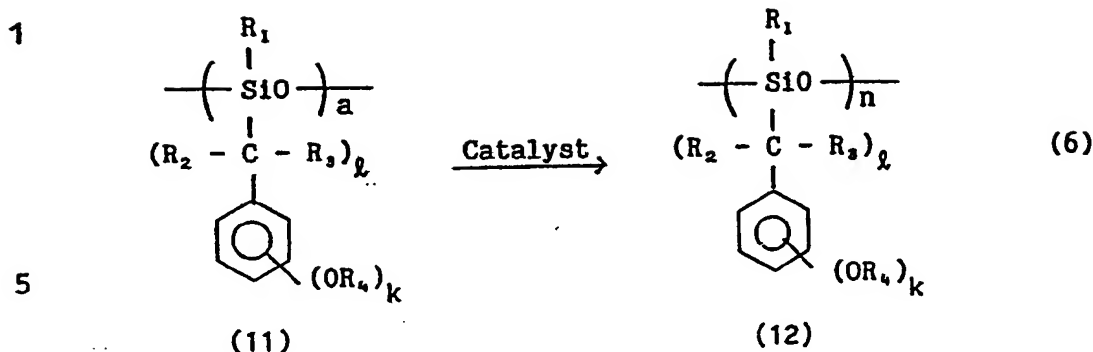


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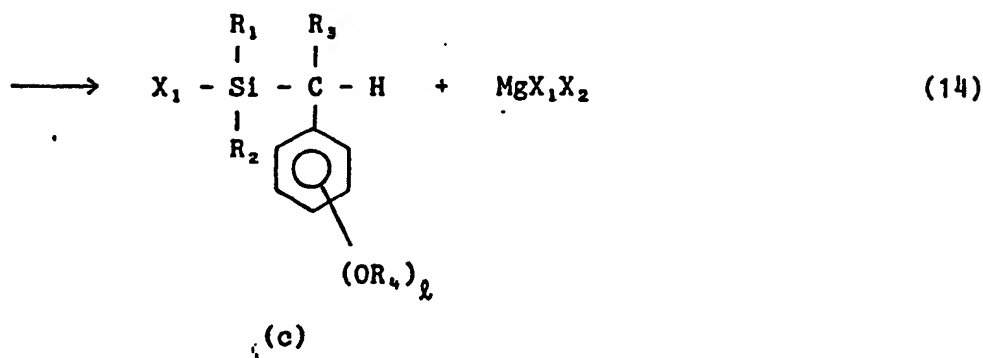
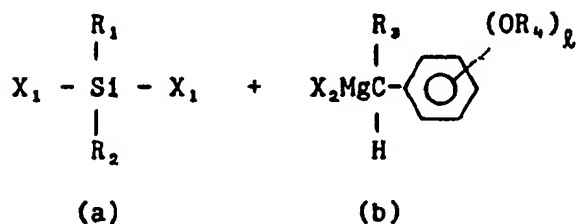


(where X_1 is a halogen (e.g., -Cl and -Br) or alkoxy group (e.g., -OMe and -OEt); X_2 is a halogen (e.g., -Cl and -Br) or alkyl group (e.g., $C_1 - C_5$ alkyl group); a is 3 to 5; and n is an integer of 3 to 1000.)

At first, a trihalogenosilane or trialkoxysilane (8) is reacted with a Grignard reagent (9) to give a dihalogenosilane or dialkoxysilane (10), according to the equation (4). Then, the dialkoxysilane (10) is hydrolyzed to give a low-molecular weight cyclic siloxane (11), according to the equation (5). The low-molecular weight cyclic siloxane (11) undergoes ring opening polymerization by the aid of a catalyst, according to the equation (6). The suitable catalyst is Me_4NOH , Bu_4POH , metal

1 hydroxide, or alkali metal-graphite interlaminar compound. The low-
 molecular weight cyclic siloxane (11) may be copolymerized with another
 kind of low-molecular weight cyclic siloxane. Finally, the OR_4 group is
 substituted by the OH group, according to the equation (7). The
 5 cleavage of the protective group may be accomplished by using $MeSiI/MeOH$
 and other reactions. Alternatively, the OH group may be protected from
 beginning by a silyl group such as t-butyldimethylsilyl group having a
 great steric hindrance and it is removed by the aid of an acid in the
 last stage.

10 The polymer (2) of this invention is usually synthesized according
 to the following equations (14) - (17).





At first, a dihalogenosilane or dialkoxysilane (a) is reacted with a Grignard reagent (b) to give a monohalogenosilane or monoalkoxysilane (c), according to the equation (14). Then, the benzyl hydrogen in the

1 compound (c) is halogenized to give the compound (d), according to the
equation (15). This reaction may be accomplished by the aid of photo-
chlorination, N-chlorosuccinimide, or N-bromosuccinimide. The compound
(d) is polymerized into the compound (e) by the aid of a metal, accord-
5 ing to the equation (16). The suitable metal is magnesium or an alkali
metal. Where magnesium is used, Si and C in the polymer skeleton are
arranged alternately; and where an alkali metal is used, the polymer
skeleton contains the Si-Si linkage and the C-C linkage. Therefore, the
polymer skeleton should be understood to contain both the Si-Si linkage
10 and C-C linkage. Finally, the conversion of the OR_n group is performed
according to the equation (17). The cleavage of the protective group
may be accomplished by using MeSiH/MeOH and other reactions. Alterna-
tively, the OH group may be protected from beginning by a silyl group
such as t-butyldimethylsilyl group having a great steric hindrance and
15 it is removed by the aid of an acid in the last stage.

The polymer (3) of this invention is produced in two steps. In the
first step, a polyorganosilsesquioxane, with the hydroxyl group
protected, is synthesized from a corresponding trichlorosilane or
trialkoxysilane. In the second step, the protective group is removed.
20 The trichlorosilane or trialkoxysilane may be produced in many ways.
Examples include the Grignard reaction, the condensation of a halide
(such as benzyl chloride derivative) and HSiCl₃, by the aid of a tertiary
amine and copper salt, and the addition of HSiCl₃ to a styrene deriva-
tive by the aid of a platinum catalyst.

1 The polyorganosilsesquioxane with the hydroxyl group protected was
synthesized according to the processes, with or without modification,
disclosed in J. Am. Chem. Soc., 82, 6194 (1960), J. Am. Chem. Soc., 87,
4317 (1965), Japanese Patent Publication No. 15989/1965, and Japanese
5 Patent Laid-open Nos. 88099/1978 and 66422/1984. The resulting polymers
had a degree of polymerization (n) in the range of 8 to 200 regardless
of the process. Therefore, the process for synthesis is not specifi-
cally limited in this invention. On the other hand, there are various
methods for removing the protective group. For example, trimethylsilyl
10 iodide is used for an alkoxy group; tetra-n-butylammonium fluoride is
used for a t-butyldimethylsiloxy group; and phosphorus pentachloride is
used for methylene acetal. All of these methods can be applied to the
synthesis of the polymer of this invention. Therefore, there are no
limits in the methods of removing the protective group in the synthesis
15 of the polymer of this invention.

The polymer of this invention is soluble in alkaline water such as
aqueous solutions of tetramethylammonium hydroxide and potassium hydrox-
ide; and it is also readily soluble in widely used organic solvents such
as alcohols, ethers, amides, ketones, esters, and cellosolves. Thus,
20 the polymer can be made into film by the aid of one of these solvents.

The polymer of this invention can be used as a base material of
photoresists for light or radiation, if it is incorporated with a photo-
sensitive or radiation-sensitive dissolution inhibitor suitable for the
light or radiation to be used. The thus produced photoresist can be
25 developed with an alkali.

1 In addition, as in the case of conventional resist of alkali development type, the polymer of this invention can be used as a positive photoresist if it is combined with o-quinonediazide, and it can also be used as a negative photoresist if it is combined with bisazide.

5 The polymer of this invention as such can be used as a negative electron resist. On the other hand, when combined with poly(olefinsulfone), it can be used as a positive electron resist.

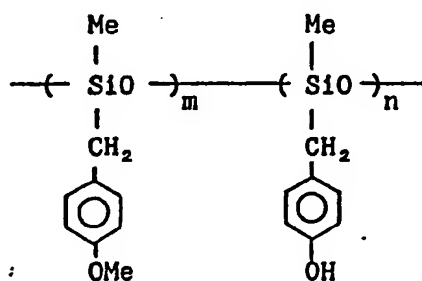
The film formed from the polymer of this invention does not waste at all in an oxygen plasma, keeping an extremely high resistance to dry
10 etching. Therefore, it can be used as the top layer resist where the double-layer resist method is employed.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

The invention is now described in more detail with reference to the following examples.

EXAMPLE 1

15 Poly(methyl-p-methoxybenzylsiloxane-co-methyl-p-hydroxybenzylsiloxane):



(a) Synthesis of methyldichloro-p-methoxybenzylsilane

In a 1-liter three-neck flask equipped with a stirrer, reflux
25 condenser, dropping funnel, and thermometer were placed 23.3 g (1.00

1 mol) of magnesium powder, 82.6 g (0.640 mol) of methyltrichlorosilane,
and 200 ml of diethyl ether. The magnesium was activated with iodine
and dibromoethane. The flask was cooled to 10°C to 0°C. A mixture of
50.0 g (0.319 mol) of p-methoxybenzyl chloride and 300 ml of diethyl
5 ether was added dropwise with stirring from the dropping funnel over 4
hours at 10°C to 0°C. After aging at room temperature for 1 hour,
excess magnesium and magnesium chloride were removed by suction
filtration. The filtrate was distilled to give 48.8 g (0.208 mol) of
the desired product in a yield of 65%.

10 The properties of the desired product were as follows:

Boiling point: 86 - 88°C/2 mmHg

NMR : (60 MHz, CCl₄, CH₂Cl₂, δ 5.33) δ 0.79 (3H, s), δ 2.68 (2H, s),
δ 3.84 (3H, s), δ 6.80 (2H, d, J=9 Hz), δ 7.18 (2H, d, J=9 Hz).

IR (ν cm⁻¹) 1620, 1520, 1480, 1310, 1260, 1190, 1100, 1050, 850, 810,

15 755.

(b) Synthesis of trimer of methyl-p-methoxybenzylsiloxane

In a 1-liter three-neck flask equipped with a stirrer, reflux
condenser, and dropping funnel were placed 200 ml of water. A mixture
of 85.6 g (0.364 mol) of p-methyldichloro-p-methoxybenzylsilane and 200
20 ml of ethyl ether was added dropwise with stirring from the dropping
funnel over 4 hours at room temperature. After aging at room tempera-
ture for 1 hour, the organic compounds were extracted with diethyl ether
and the extract was washed with water until the washings became neutral.
The diethyl ether solution was dried with sodium sulfate and diethyl

25

1 ether was distilled away. Distillation with calcium hydride gave 20.3 g (37.5 mmol) of the desired product in a yield of 31%.

The properties of the desired product were as follows:

Boiling point: 241°C/0.8 mmHg to 253°C/0.13 mmHg

5 NMR : (60 MHz, CCl₄, CH₂Cl₂, δ 5.33) δ 0.02, δ 0.03, δ 0.16 (9H, sx3), δ 1.96, 2.07 (6H, sx2), δ 3.79 (9H, s), δ 6.74 (CH, d, J=9 Hz), δ 6.98 (6H, d, J=9 Hz).

IR (ν cm⁻¹) 1620, 1520, 1470, 1305, 1260, 1220, 1185, 1090, 1030, 840, 815, 770.

10 (c) Synthesis of poly(methyl-p-methoxybenzylsiloxane)

In a 100-ml eggplant-shape flask containing a magnetic bar were placed 9.0 g (17 mmol) of trimer of methyl-p-methoxybenzylsiloxane and 45 mg (0.5 wt%) of tetramethylammonium hydroxide. With the flask stoppered and immersed in an oil bath at 80°C, the reactants were stirred by a magnetic stirrer. The weight-average molecular weight reached 5,000 four minutes after the start of reaction. It reached 10,000 after 10 minutes, 14,000 after 20 minutes, 16,000 after 30 minutes, and 19,000 after 60 minutes. The desired product having a desired weight-average molecular weight can be obtained when the flask is removed from the oil bath and cooled with water after a certain period of reaction time. The content in the flask was used as such for the subsequent reaction.

The properties of the desired product were as follows:

NMR : (60 MHz, CCl₄, CH₂Cl₂, δ 5.33) δ 0.00 (3H, br. s), δ 1.94 (2H, br. s), δ 3.71, δ 3.80 (3H, sx2), δ 6.82 (4H, br. s).

1 IR (ν cm^{-1}) 2975, 2850, 1620, 1520, 1475, 1305, 1260, 1225, 1190, 1090,
1040, 845, 810, 775, 755.

(d) Synthesis of poly(methyl-p-methoxybenzylsiloxane-co-methyl-
p-hydroxybenzylsiloxane)

5 Poly(methyl-p-methoxybenzylsiloxane) was placed in a flask, and
flask was equipped with a reflux condenser and the atmosphere in the
flask was replaced with nitrogen. 30 ml of chloroform was added to the
flask to make a uniform solution of the polymer. A prescribed amount of
trimethylsilyl iodide was added. After stirring at room temperature for
10 3 hours, 20 ml of methanol was added and stirring was continued for 2
hours. After the completion of the reaction, the low-boiling fractions
were distilled away at room temperature under reduced pressure. The
residues were extracted with tetrahydrofuran and diethyl ether. The
extract was washed with an aqueous solution of sodium hydrogen sulfite,
15 an aqueous solution of sodium hydrogen carbonate, and finally an aqueous
solution of sodium chloride. The solvent was distilled away under
reduced pressure. Thus there was obtained the desired product.
Trimethylsilyl iodide in an amount of 1.5 equivalent for the methoxy
group converted 100% of methoxy groups into hydroxyl groups. An amount
20 of 1.3 equivalent resulted in a 90% conversion, 0.76 equivalent, 71%,
0.65 equivalent, 58%, and 0.47 equivalent, 41%.

The properties of the desired product were as follows:

Melting point: (polymer having a weight-average molecular weight of
10,000 and 80% converted hydroxyl groups) 53 - 68°C

- 1 NMR : (60 MHz, CD_3SCD_3 , CH_2Cl_2 , δ 5.68) δ 0.07 (br. s, CH_3), δ 1.85 (br. s, $-\text{CH}_2-$), δ 4.00 (br. s, $-\text{OCH}_3$), δ 6.61 (br. s, ring proton), δ 8.81 (br. s, $-\text{OH}$).
IR ($\nu \text{ cm}^{-1}$) 3320, 1620, 1510, 1450, 1270, 1240, 1180, 1070, 840, 800.

5 (e) Solubility

Poly(methyl-p-methoxybenzylsiloxane-co-methyl-p-hydroxybenzylsiloxane) was examined for solubility toward commonly used organic solvents. It was found to be soluble in methanol, tetrahydrofuran, N,N-dimethylacetamide, 2-methylcyclohexanone, isoamyl acetate, methyl
10 cellosolve, and dimethylsulfoxide, and also found to be insoluble in toluene, hexane, and carbon tetrachloride.

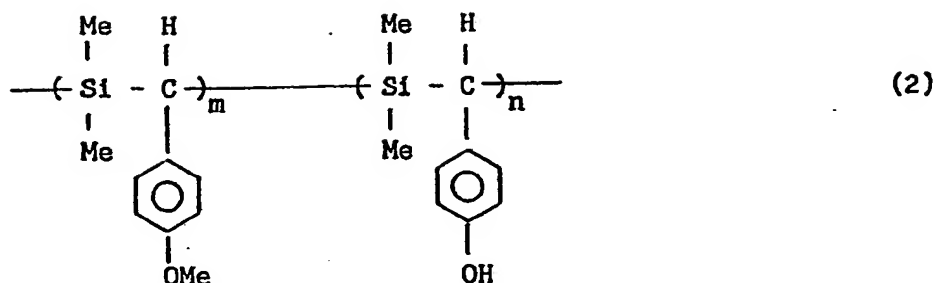
It was found to be soluble in aqueous solutions of tetramethyl ammonium hydroxide and potassium hydroxide.

(f) Resistance to oxygen plasmas

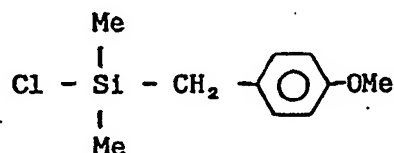
- 15 A 5 wt% solution of poly(methyl-p-methoxybenzylsiloxane-co-methyl-p-hydroxybenzylsiloxane) in 2-methylcyclohexanone was applied to a silicon substrate by spin coating method. A 0.2 μm thick polymer film was formed. The film was baked at 90°C for 30 minutes. The baked film was exposed to oxygen plasmas (oxygen pressure: 0.5 Torr, degree of
20 vacuum: 10^{-4} Torr, RF: 300 W) for 10 minutes. The film did not waste at all.

EXAMPLE 2

Poly(1,1-dimethyl-2-p-methoxyphenylsilylmethylene-co-1,1-dimethyl-2-p-hydroxyphenylmethylene)



(a) Synthesis of chlorodimethyl-p-methoxybenzylsilane



In a 2-liter three-neck flask equipped with a stirrer, reflux condenser, dropping funnel, and thermometer were placed 31.0 g (1.28 mol) of magnesium powder, 129 g (1 mol) of dimethyldichlorosilane, and 500 ml of diethyl ether. With the flask cooled below 10°C, a mixture of 100 g (0.639 mol) of p-methoxybenzyl chloride and 250 ml of diethyl ether was added dropwise from the dropping funnel over 4 hours. After aging at room temperature for 1 hour, excess magnesium and magnesium chloride were removed by suction filtration. The filtrate was distilled to give 87.8 g (0.409 mol) of the desired product in a 64.0% yield.

The properties of the desired product were as follows:

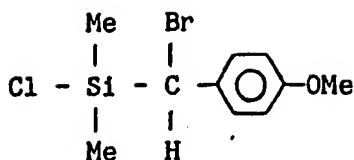
Boiling point: 80°C/1 mmHg

NMR : (60 MHz, CCl₄, CH₂Cl₂, δ 5.33) δ 0.46 (6H, s), δ 2.40 (2H, s), δ 3.83 (3H, s), δ 6.82 (2H, d, J=9.0 Hz), δ 7.08 (2H, d, J=9.0 Hz).

IR (ν cm⁻¹) 2970, 1620, 1520, 1475, 1310, 1260, 1195, 1090, 1055, 855, 825, 780.

(b) Synthesis of bromo-p-methoxyphenylchlorodimethylsilylmethane (dl)

1



In a 500-ml three-neck flask equipped with a stirrer and reflux
5 condenser were placed 43.2 g (0.200 mol) of chlorodimethyl-p-methoxy-
benzylsilane and 200 ml of carbon tetrachloride. With heating and
stirring, 35.6 g (0.200 mol) of N-bromosuccinimide was added slowly.
After aging for 1 hour, succinimide was removed by suction filtration
and the filtrate was distilled to give 31.7 g (0.108 mol) of the desired
10 product in a yield of 54.0%.

The properties of the desired product were as follows:

Boiling point: 113 - 117°C/2.5 mmHg

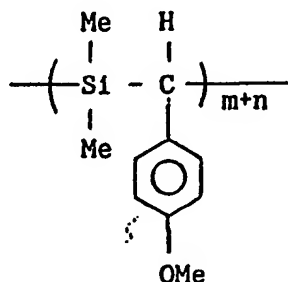
Melting point: lower than 30°C.

NMR : (60 MHz, CCl_4 , TMS) δ 0.50 (3H, s), δ 0.60 (3H, s), δ 3.73
15 (3H, s), δ 4.32 (1H, s), δ 6.73 (2H, d, J=9 Hz), δ 7.19 (2H, d, J=9 Hz).

IR (ν cm^{-1}) 1610, 1510, 1465, 1305, 1260, 1185, 1040, 865, 845, 820,
800.

(c) Synthesis of poly(1,1-dimethyl-2-p-methoxyphenylmethylene)

20



In a 500-ml three-neck flask equipped with a Henschberg stirrer,
25 reflux condenser, and dropping funnel, with the atmosphere therein

1 replaced with nitrogen, were placed 7.6 g (0.33 gram atom) of sodium and
120 ml of toluene. The contents were heated with stirring to give fine
sodium particles. Under the refluxing of toluene with stirring, a
mixture of 44.1 g (0.15 mol) of bromo-p-methoxyphenylchlorodimethyl-
5 silylmethane and 30 ml of toluene was added dropwise over 1 hour. After
aging for 1 hour, the reaction product was poured into methanol to
removed excess sodium and precipitate the polymer. After filtration,
the polymer was dissolved in tetrahydrofuran and the resulting solution
was added dropwise to water to remove the sodium salt. The filtered
10 polymer was washed with methanol and dried under reduced pressure to
give 10 g of the desired product in a yield of 37%.

The properties of the desired product were as follows:

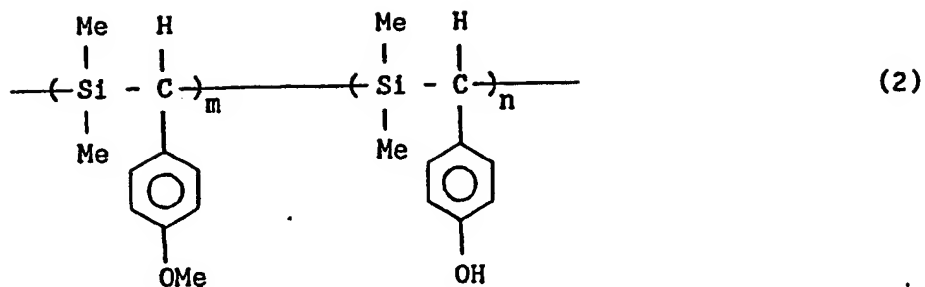
Softening point: 121 - 125°C

Weight-average molecular weight: 3,400

15 NMR : (60 MHz, C_6D_6 , CH_2Cl_2 , δ 5.33) δ 1.27 (6H, br. s), δ 4.43 (3H,
br. s), δ 7.4 - 8.7 (4H, br. s). (Methyl proton was not confirmed.)
IR (ν cm^{-1}) 2960, 2850, 1610, 1510, 1470, 1385, 1300, 1260, 1100, 1045,
860, 790.

The polymer formed through Grignard reaction had a weight-average
20 molecular weight of about 1000.

(d) Synthesis of poly(1,1-dimethyl-2-p-methoxyphenylmethylene-co-1,1-
dimethyl-2-p-hydroxyphenylsilmethylene)



In a 25-ml eggplant-shape flask equipped with a reflux condenser were placed 0.50 g (2.8 mmol in terms of monomer units) of poly(1,1-dimethyl-2-p-methoxyphenylsilmethylene) having a weight-average molecular weight of 3,000, 1.5 ml of chloroform, and 0.88 g (4.4 mmol) of trimethylsilyl iodide. Reaction was carried out for 4 hours with stirring by a magnetic bar. 4 ml of methanol was added and stirring was continued for 3 hours. Low boiling fractions were distilled away under reduced pressure at room temperature, and the residues were extracted with ether. The ether solution was washed with an aqueous solution of sodium hydrogen sulfite, an aqueous solution of sodium hydrogen carbonate, and finally an aqueous solution of sodium chloride. Ether was distilled away under reduced pressure. The resulting polymer was reprecipitated from tetrahydrofuran/water. After thermodrying under reduced pressure, there was obtained 0.23 g of the desired product.

20 The properties of the desired product were as follows:

Weight-average molecular weight: 1,800

OH content: 100%

Softening point: 107 - 113°C

NMR : (60 MHz, DMSO-d₆, δ 5.68) δ 0.79-0.51 (6H, br. s), 6.15-7.35 (4H,

25 br. s). Methyl proton and hydroxy proton were not confirmed.

1 IR (ν cm^{-1}) 3380, 2980, 1510, 1270, 1100, 1040, 860, 810.

In the meantime, the OH content can be controlled by the amount of trimethylsilyl iodide or the reaction time. For example, 90% of methoxy groups is converted into hydroxyl groups where the reaction time is 4
5 hours and the amount of trimethylsilyl iodide is 1.3 equivalent for methoxy groups. 0.80 equivalent gave a conversion of 70%, 0.65 equivalent, 55%, and 0.50 equivalent, 40%.

The OH content was determined by performing the reaction in deuteriochloroform. During the reaction, the conversion of methoxy groups into
10 trimethylsiloxy groups was traced by means of NMR spectrometry.

(e) Solubility

Poly(1,1-dimethyl-2-p-methoxyphenylsilmethylene-co-1,1-dimethyl-2-p-hydroxyphenylsilmethylene) was examined for solubility toward commonly used organic solvents. The polymer containing more than 40% of hydroxyl
15 groups was found to be soluble in methanol, tetrahydrofuran, N,N-dimethylacetamide, 2-methylcyclohexanone, isoamyl acetate, methyl cellosolve, and dimethylsulfoxide, and also found to be insoluble in toluene, hexane, and carbon tetrachloride.

It was found to be soluble in aqueous solutions of tetramethyl
20 ammonium hydroxide and potassium hydroxide.

(f) Resistance to oxygen plasmas

A 10 wt% solution of poly(1,1-dimethyl-2-p-methoxyphenylsilmethylene-co-1,1-dimethyl-2-p-hydroxyphenylsilmethylene) in 2-methylcyclohexanone was applied to a silicon substrate by spin coating method.
25 Upon baking at 100°C for 30 minutes, a 0.2 μm thick polymer film was

- 1 formed. The baked film was exposed to oxygen plasmas (oxygen pressure: 0.5 Torr, degree of vacuum: 10^{-4} Torr, RF: 300 W) for 10 minutes. The film did not waste at all.

EXAMPLE 3

- 5 Poly(hydroxybenzylsilsesquioxane-co-p-methoxybenzylsilsesquioxane):

(a) Synthesis of p-methoxybenzyltrichlorosilane

- In a 2-liter three-neck flask equipped with a stirrer, reflux condenser, dropping funnel, and thermometer were placed 30 g (1.2 gram atom) of magnesium powder, 170 g (1.00 mol) of silicon tetrachloride, 10 and 500 ml of diethyl ether. The flask was cooled below 10°C . A mixture of 100.0 g (0.639 mol) of p-methoxybenzyl chloride and 200 ml of diethyl ether was added dropwise with stirring from the dropping funnel over 4 hours. After aging at room temperature for 1 hour, excess magnesium and magnesium chloride were removed by suction filtration.
- 15 The filtrate was distilled to give 44.0 g (0.172 mol) of the desired product in a yield of 26.9%.

The properties of the desired product were as follows:

Boiling point: $117.5 - 119.5^{\circ}\text{C}/3.0 \text{ mmHg}$

- NMR : (60 MHz, CCl_4 , CH_2Cl_2 , δ 5.33) δ 2.91 (2H, s), δ 3.90 (3H, s), 20 δ 6.91 (2H, d, $J=8 \text{ Hz}$), δ 7.20 (2H, d, $J=8 \text{ Hz}$).

(b) Synthesis of poly(p-methoxybenzylsilsesquioxane)

- In a 100-ml three-neck flask equipped with a magnetic bar, dropping funnel, and reflux condenser were placed 11 g (0.13 mol) of sodium hydrogen carbonate and 40 ml of water. A mixture of 10.23 g (40.00 25 mmol) of p-methoxybenzyltrichlorosilane and 10 ml of diethyl ether was

1 added dropwise from the dropping funnel over 30 minutes, followed by
aging for 30 minutes. After the completion of the reaction, the
reaction mixture was extracted with ether, and the extract was dried
with sodium sulfate. Diethyl ether was distilled away under reduced
5 pressure to give 5.10 g of hydrolyzate.

The properties of the hydrolyzate were as follows:

NMR : (60 MHz, CDCl_3 , CH_2Cl_2 , δ 5.33) δ 2.03 (2H, br. s), δ 3.80 (3H,
br. s), δ 6.80 (4H, br. s).

IR ($\nu \text{ cm}^{-1}$) 3400, 2950, 2850, 1610, 1510, 1460, 1300, 1250, 1180, 1090,
10 1035, 890, 835, 790, 760.

Weight-average molecular weight: 2,000

In a 25-ml eggplant-shape flask were placed 4.80 g of the
hydrolyzate obtained as mentioned above and 49 mg of 10 wt% solution of
potassium hydroxide in methanol. The contents were heated at 200°C for
15 2 hours. After the completion of the reaction, the reaction mixture was
dissolved in benzene and the benzene solution was added dropwise to
methanol to precipitate solids. The solids were filtered off and dried
to give 4.00 g of poly(p-methoxybenzylsilsesquioxane).

NMR : (60 MHz, CDCl_3 , CH_2Cl_2 , δ 5.33) δ 1.19 (2H, br. s), δ 3.78 (3H,
20 br. s), δ 6.73 (4H, br. s).

IR ($\nu \text{ cm}^{-1}$) 2950, 2850, 1615, 1515, 1465, 1305, 1250, 1195, 1120, 1040,
840, 800, 770.

Weight-average molecular weight: 3,300

(c) Synthesis of poly(hydroxybenzylsilsesquioxane-co-p-methoxybenzyl-
25 silsesquioxane):

1 In a 100-ml eggplant-shape flask equipped with a reflux condenser
were placed 3.73 g (21.6 mmol in terms of $\text{MeOC}_6\text{H}_4\text{CH}_2\text{SiO}_{1/2}$), 20 ml of
chloroform, and 6.92 g (34.6 mmol) of trimethylsilyl iodide. Reaction
was carried out at 70°C for 22 hours with stirring by a magnetic bar.
5 20 ml of methanol was added at room temperature and stirring was
continued for 30 minutes. Low boiling fractions were distilled away
under reduced pressure, and the residues were extracted with a mixed
solvent of diethyl ether and tetrahydrofuran. The extract was washed
10 with an aqueous solution of sodium hydrogen sulfite, an aqueous solution
of sodium hydrogen carbonate, and finally an aqueous solution of sodium
chloride. The solvent was distilled away under reduced pressure. The
resulting polymer was reprecipitated from acetone/hexane. After
thermodrying under reduced pressure, there was obtained 2.71 g of the
desired product.

15 The properties of the desired product were as follows:

Weight-average molecular weight: 4,800

OH content: 85%

NMR : (60 MHz, DMSO- d_6 , δ 5.68) δ 1.75 (2H, br. s), 3.63 (-OCH₃, br. s),
6.58 (4H, br. s), 8.88 (-OH, br. s).

20 IR (ν cm⁻¹) 3350, 1620, 1515, 1450, 1240, 1185, 1120, 1040, 840, 805,
760.

In the meantime, the OH content can be controlled by the amount of
trimethylsilyl iodide or the reaction time. For example, where 1.6
equivalent of trimethylsilyl iodide was used, the reaction time of 4
25 hours brought about a 36% conversion of methoxy groups into hydroxyl

1 groups, 7 hours, 54%, 12 hours, 75%, 48 hours, 95%, and 72 hours, almost
100%. The OH content was determined by performing the reaction in
deuteriochloroform. During the reaction, the conversion of methoxy
groups into trimethylsiloxy groups was traced by means of NMR
5 spectrometry.

(d) Solubility

Polymer of this invention was examined for solubility toward
commonly used organic solvents. The polymer having more than 50% of
hydroxyl groups were soluble in methanol, tetrahydrofuran, N,N-dimethyl-
10 acetamide, 2-methylcyclohexanone, isoamyl acetate, ethyl cellosolve, and
dimethylsulfoxide, and were insoluble in toluene, hexane, and carbon
tetrachloride.

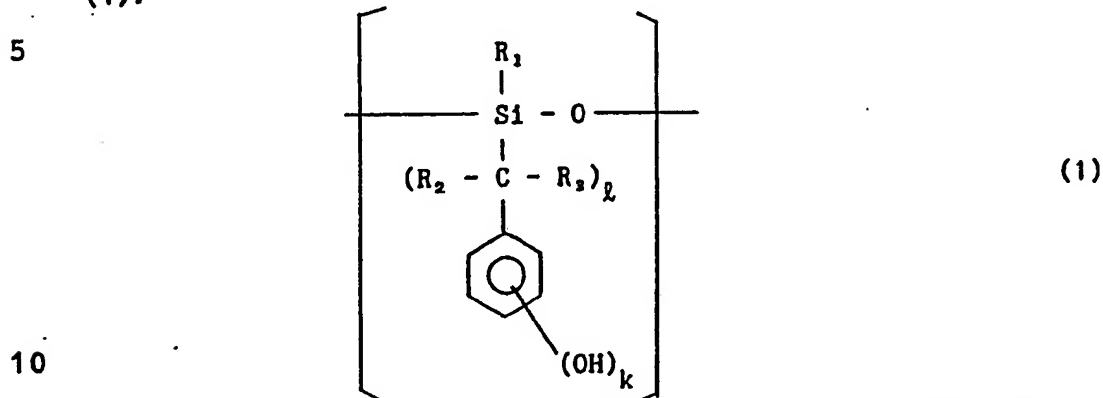
It was also soluble in an aqueous solution of tetramethyl ammonium
hydroxide.

15 (e) Resistance to oxygen plasmas

An 8 wt% solution of the polymer of this invention in 2-methyl-
cyclohexanone was applied to a silicon substrate by spin coating method.
Upon baking at 100°C for 30 minutes, there was obtained a 0.2 μ m thick
film. The baked film was exposed to oxygen plasmas (oxygen pressure:
20 0.5 Torr, RF: 300 W, barrel type asher) for 20 minutes. The film did
not waste at all.

1 WHAT IS CLAIMED IS:

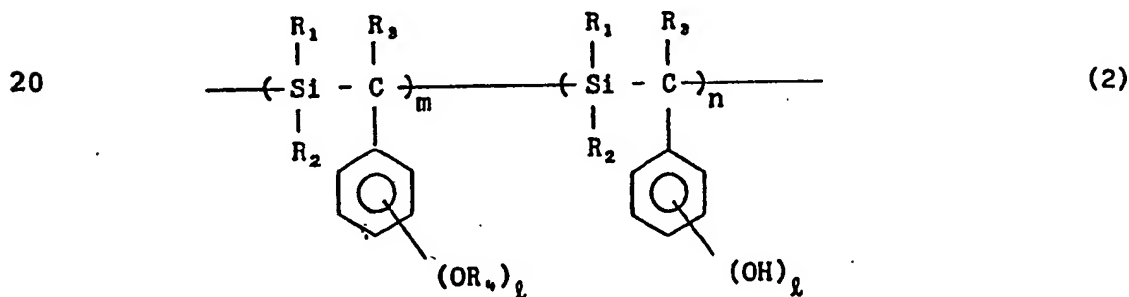
1. An alkali-soluble siloxane polymer in which more than 40% of the siloxane skeleton is composed of the unit represented by the formula (1).



(where R_1 is hydrogen or a monovalent organic group other than aromatic groups; R_2 and R_3 are each hydrogen, chlorine, or a monovalent organic group; l is an integer of 1 to 6; and k is 1 or 2.)

2. An alkali-soluble siloxane polymer as set forth in Claim 1, wherein R_1 is methyl group, R_2 and R_3 are each hydrogen, l is 1, k is 1, and the OH group is at the p-position of the benzene ring.

3. an alkali-soluble silmethylen polymer composed of the constitutional unit represented by the formula (2).



(where R_1 , R_2 , and R_3 are each a monovalent organic group; R_4 is an alkyl group; l is 1 or 2; the OH groups and OR_4 groups are at arbitrary

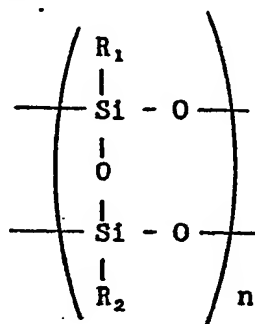
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1 positions on the benzene ring, m and n are positive integers and m+n is
2 to 200 and n/(m+n) is greater than 0.4.)

4. An alkali-soluble silmethylen polymer as set forth in Claim 3,
wherein R₁, R₂, and R₄ are methyl groups, R₃ is hydrogen, l is 1, and
5 the OH group and OR₄ group are each on the p-position of the benzene
ring.

5. An alkali-soluble polyorganosilsesquioxane polymer represented
by the formula (3).

10



(3)

(where R₁ and R₂ are the same or different organic groups; and n is a
15 degree of polymerization), said polymer containing more than 40% of R₁
and R₂ each having phenolic hydroxyl groups.

6. An alkali-soluble polyorganosilsesquioxane polymer as set forth
in Claim 5, wherein R₁ and R₂ are each p-hydroxybenzyl group and
p-methoxybenzyl group, and n is 8 to 200.

20

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